EC CERTIFICATE

EC QUALITY SYSTEM (MODULE D) CERTIFICATE
No. 2821-MED-0002

This is to certify that: UL International (Netherlands) B.V. did undertake the relevant quality assurance assessment procedures for the quality system (Module D) of the manufacturer identified below which was found to be in compliance with the requirements of Marine Equipment Directive (MED) 2014/90/EC, last amended by Regulation EU 2020/1170, for the type described in the EC Type Examination Certificates (Module B) as listed in the Annex to this Certificate, subject to any conditions attached hereto.

Manufacturer:	**Datrex Inc.**
Address:	**PO Box 1537, 13878 Hwy 165, Kinder, LA 70648. US**
Authorised Representative:	**Mr. Lars Lund**
Address:	**Svanbervägen 9, S-69141 Karlskoga, Sweden.**

Scope: (MED Item designation)

MED/1.1 **Lifebuoys**

The attached (*schedule of approval*) forms part of this certificate.

This certificate remains valid unless cancelled, expired or revoked.

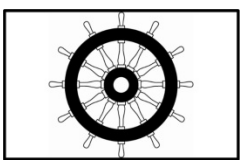
Date of re-issue:	**12 February 2021**	Issued by:	**UL International (Netherlands) B.V.**
Issue	**1**		**Notified Body 2821**
Expiry date:	**28 February 2025**		
		Signed:	
This Certificate consists of 2 pages		Name:	Horst Thelen Head of Notified Body

Notes

1: This certificate authorises the manufacturer or his authorised representative established within the Community in conjunction with the EC TYPE EXAMINATION (MODULE B) CERTIFICATE of the equipment listed in the scope to affix the “Mark of Conformity” (wheelmark).

2: This certificate loses its validity if the manufacturer makes any changes or modifications to the approved quality system, which have not been notified to, and agreed with the notified body named on this certificate and/or after lapse of time, withdrawal or revocation of the EC TYPE EXAMINATION (MODULE B) CERTIFICATE.

3: Example for the Application of the “Mark of Conformity”:



2821 Number of the Notified Body responsible for quality surveillance module
YYYY Year in which the mark is affixed

UL International (Netherlands) B.V.
Westervoortsedijk 60,
6827AT Arnhem, The Netherlands

EC CERTIFICATE

Schedule of Approval (Annex)
No. 2821-MED-0002

Date of re-issue:	**12 February 2021**	Issued by:	**UL International (Netherlands) B.V.**
Issue	**1**		**Notified Body 2821**
Expiry date:	**28 February 2025**		

Places of production:

Datrex, Inc.
PO Box 1537, 13878 Hwy 165, Kinder, LA 70648. US.

Product details:
The following is a non-exhaustive list of MED products, Technical Files, and/or EU type-examination certificates which were subject of the quality assurance assessment procedure noted below in 'Approval documentation' when the certificate was first established. Module B certificates issued subsequent to this certificate shall be covered by this certificate should they fall within the Scope as detailed on page 1, and are produced at those addresses listed above.

Module B Certificate No.	Technical File Ref.	Product Code(s)
2821-MED-0001	DatMedTechDoc,	DX0325D
2821-MED-0006	Revision 4, Issued	DX0340D
	July 13, 2017; Revised	
	Sept. 10 2020.	

Approval documentation:

UL-CSA-DatrexInc-10-2017
UL-CSA-DatrexInc-10-2018
UL-CSA-DatrexInc-11-2019
UL-JS-DatrexInc-01-2021

Limitations on the validity of certification:

None

Terms and Conditions:

1. This certificate remains the property of UL International (NL) B.V., herein "UL NL", and will be withdrawn if any conditions attached to its issue are not complied with.
2. This certificate is issued subject to the Global Services Agreement (GSA) and MED Service Terms.
3. Production is limited to the site(s) as listed, or detailed within the Technical documentation held by UL Netherlands.
4. Any system change, production/process changes, or changes in state of the art which may affect conformity shall be notified to UL Netherlands.
5. This certificate authorizes the use of the Mark of Conformity (the 'Wheel mark'), and should be referenced within the Manufacturer's Declaration of Conformity prior to placement on the market.
6. The certificate remains valid provided the annual audit schedule is maintained and the conclusions obtained attest to continued conformity of the product(s) and quality-control system with the Marine Equipment Directive 2014/90/EU requirements.

UL International (Netherlands) B.V.
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6827AT Arnhem, The Netherlands